

PCTWORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁷ : C07K 14/01, 16/08, G01N 33/68, A61P 31/20	A1	(11) International Publication Number: WO 00/66621 (43) International Publication Date: 9 November 2000 (09.11.00)
(21) International Application Number: PCT/EP00/03958 (22) International Filing Date: 3 May 2000 (03.05.00) (30) Priority Data: 9901601-6 4 May 1999 (04.05.99) SE (71) Applicant (for all designated States except US): TRIPEP AB [SE/SE]; Hålsövägen 7, S-141 57 Huddinge (SE). (72) Inventor; and (75) Inventor/Applicant (for US only): SÄLLBERG, Matti [SE/SE]; Stora Rönnens Gränd 15, S-181 68 Täby (SE). (74) Agents: NILSSON, Brita et al.; AB Stockholms Patentbyrå, Zacco & Bruhn, P.O. Box 23101, S-104 35 Stockholm (SE).		(81) Designated States: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>
(54) Title: PEPTIDES FROM THE TT VIRUS SEQUENCE AND MONOSPECIFIC ANTIBODIES BINDING TO THE TT VIRUS (57) Abstract The peptide having the amino acid sequence SEQ ID NO:1, and optionally this peptide in mixture with one or more peptides SEQ ID NO:3 - 11, is described. All these peptides correspond to regions of the genomic TT virus sequence. Further, monospecific antibodies binding to the TT virus are disclosed. The peptides may be coupled to a carrier and/or label, or immobilized on a solid phase. The peptide or peptide mixture, or the monospecific antibody may be used in a medicament or in diagnostic kits. The peptide or peptide mixture may also be used for immunization of a non-human mammal to produce monospecific antibodies directed against TT virus.		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakhstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

Peptides from the TT virus sequence and monospecific antibodies binding to the TT virus.

The present invention relates to peptides derived from the genomic TT virus sequence and monospecific antibodies binding to the TT virus. Further, the invention relates to the peptides and antibodies of the invention for respective use in medicaments. Diagnostic kits comprising the peptides of the invention as diagnostic antigens, and diagnostic kits comprising the antibodies of the invention as diagnostic antigens are also comprised by the invention. The peptides of the invention may be used for immunization of a non-human mammal to produce monospecific antibodies directed against TT virus.

Background

In 1997 a novel human infectious agent was identified from the serum of a Japanese patient with post transfusion non A-G hepatitis and named TT virus (TTV) [1]. TTV DNA was detected in 47% of patients with fulminate non-A-G hepatitis and 46% of patients with chronic liver disease of unknown etiology [2] suggesting that TTV may be the cause of some idiopathic liver disease. TTV is global [3]] and seems to be more common in populations with increased risk for infection with blood borne viruses [2] e.g. hemophiliacs and drug addicts. However, non-parenteral transmission seems also to be possible [2].

TTV is a non-enveloped, single stranded DNA virus with a genome of at least 3,7 kb [4]. It has a range of sequence divergence, allowing classification into different genotypes and subtypes [4]. A relationship with the family *Parvoviridae* has been discussed [4]. Subsequent analyses revealed evidence of hepatotropism of TTV [2] and in some patients with non A-G post transfusion hepatitis and TTV viremia TTV DNA titres correlated with aminotransferase levels [1].

However, an evidence for an association between TTV infection and severe liver disease could not be strengthened [3, 5, 6]. The epidemiological, immunological, and clinical significances of TTV infections are still uncertain. Moreover, no serological tests for TTV infection are available yet and at the moment PCR is the only available diagnostic tool.

It would be desirable to be able to diagnose TTV infection in man, and to develop medicaments based on peptides for immunization and/or antibodies against TTV.

Description of the invention

The present invention is based on synthetic peptides that correspond to different regions of the genomic sequence from the recently described TT virus (TTV; [1]). A total of 80 overlapping peptides corresponding to the two open reading frames (ORFs; Genbank

accession no AB008394) 1 and 2 were synthesized. These were analyzed with eight human serum samples with TTV infection and eight human samples without TTV infection. Reactive human serum samples all reacted with a peptide with the sequence SEQ ID NO:1 :

TATTTTYAYPGTNRPPV. The reactivities could be fine mapped to the sequence SEQ ID
5 NO:2 : YAYPGTNRPPV where the residues PV were found to be those most essential for the binding of human antibodies.

Thus, the present invention is directed to a peptide having the amino acid sequence
SEQ ID NO: 1

TATTTTYAYPGTNRPPV

10 wherein one to all six of the N-terminal amino acids TATTTT may be omitted.

In an embodiment of the invention the peptide has the amino acid sequence SEQ ID
NO:2
YAYPGTNRPPV.

The invention is also directed to a peptide mixture comprising the peptide SEQ ID
15 NO: 1

TATTTTYAYPGTNRPPV

wherein one to all six of the N-terminal amino acids TATTTT may be omitted, and at least
one other of the peptides listed in Table 4 having the amino acid sequences SEQ ID NO:3 -
11.

20 The peptide and/or at least one of the peptides in the peptide mixture of the invention may be coupled to a carrier and/or label. Examples of carriers are plastic surfaces, such as microplates, beads etc.; organic molecules such as biotin; proteins, such as bovine serum albumin; peptide linkers, or polypeptides. Examples of labels that can be used, primarily for diagnostic purposes, are radioactive isotopes, enzymes, fluorescent markers, etc.

25 Further, the peptide and/or at least one of the peptides in the peptide mixture of the invention may be immobilized on a solid phase, such as a glass or plastic surfaces, primarily for diagnostic purposes or purification of antibodies.

The present invention is also directed to the peptide or peptide mixture of the invention for use in a medicament, optionally coupled to or in combination with other
30 biologically active or inactive ingredients, such as a vaccine for prevention of TT virus infection.

Further, the invention is directed to monospecific antibodies binding to the TT virus.

In an embodiment of the invention, the monospecific antibody binds to an amino acid sequence selected from the group consisting of the amino acid sequences SEQ ID NO:1 - 11.

5 The invention is additionally directed to a monospecific antibody according to the invention for use in a medicament, optionally coupled to or in combination with other biologically active or inactive ingredients, such as a medicament for administration to a patient already infected with TTV.

10 The present invention is also directed to a diagnostic kit comprising a peptide or peptide mixture according to the invention as diagnostic antigen(s). The kit may be used in an immunological assay, such as EIA, RIA etc, to detect the presence of antibodies against TTV in a biological fluid, such as blood or plasma.

15 The invention is further directed to a diagnostic kit comprising one or more monospecific antibody according to the invention as diagnostic antibodies. The kit may be used in an immunological assay, such as EIA, RIA etc, to detect the presence of antibodies against TTV in a biological fluid, such as blood or plasma.

The diagnostic kits will normally comprise additional ingredients for performing an immunological assay. These additional ingredients will depend on the actual assay to be used and will often comprise positive and negative standard serum samples and written instructions for use.

20 The present invention is additionally directed to the use of a peptide according to the invention for immunization of a non-human mammal to produce monospecific antibodies directed against TT virus.

25 The present invention will now be further illustrated by reference to the following description of experiments and specific embodiments of the invention, which are not to be considered as limitations to the scope of the invention defined in the claims.

Description of experiments

Serum samples:

30 Coded serum samples were obtained from a serum bank containing healthy blood donors, children with or without liver disease, mothers with IVDU (intravenous drug use) and their children.

PCR amplification for the detection of TTV DNA in serum:

Total DNA was isolated from 50 µl patient serum by phenol/chloroform purification. The DNA of all patients was analyzed with two different primer settings by (semi) nested PCR. Five µl patient DNA were added to a 45 µl reaction mix containing 1 U

taq polymerase (Perkin-Elmer Applied Biosystems, Norwalk, CO), 10x PCR buffer, 200 μ mol $MgCl_2$, dNTPs (125 μ mol/nucleotide) and 20 pmol of each primer. The first round primers were 5TTVout5 (5'-ACA GAC AGA GGA GAA GGC AAC ATG- 3') and either 3TTVout (5'- CTG GCA TTT TAC CAT TTC CAA AGT T- 3') or 3TTXout (5'-TAC CAY TTA GCT CTC ATT CTW AT- 3') as downstream primers. The DNA was amplified as follows: 95°C for 4.5 minutes and then 33 cycles of 95°C for 30 sec, 50°C for 30 sec and 72°C for 1 min, and at the end 72°C for 4 min. A second round PCR was performed using 5 μ l of the first-round PCR product under identical conditions. The second round inner primers were either 5TTVIn (5'-GGC AAC ATG YTR TGG ATA GAC TGG - 3') or 5TTVXin (5'-ACA GGA GAC HMA AAC ATA SA- 3') as upstream primers and 3TTVout. The correct size of about 275 respectively 140 bp was determined by agarose gel electrophoresis (3%). Samples which were either positive with both primer sets or reproducibly positive with one primer set were considered as TTV positive. Primer sequences were based on Genebank accession no AB008394).

15 Peptide synthesis

Overlapping peptides (18 aa long with a 8 aa overlap) corresponding to the ORF1 and ORF2 of TTV (Table 1; Genebank accession no AB008394) were produced by a multiple peptide synthesizer using standard Fmoc chemistry [7] (Syro, Syntex, Germany).

Detection of human antibodies in serum

20 The EIAs mainly followed previous protocols [8]. Microplates (Nunc, Denmark) where coated for 48 hours with synthetic peptides at a concentration of 10 μ g/ml in 0.05M sodium carbonate buffer pH 9.6. After blocking for 2 hours at room temperature with phosphate buffered saline containing 1% bovine serum albumin, 2% goat serum and 0.05% Tween 20 (dilution buffer) the plates were incubated with human sera diluted 1:100 in
25 dilution buffer. Bound human IgG was indicated by incubation with anti-human IgG antibodies conjugated to alkaline phosphatase (Sigma Chemicals, St. Louis, MO). The plates were developed by the addition of dinitro-phenylene-diamine (Sigma) and the optical densities were determined at 405nm.

Immunization and induction of TTV-specific antibodies

30 Groups of Balb/c were immunized intra peritoneally with 100 μ g of the TTV peptide 35 (SEQ ID NO:1) emulsified 1:1 in complete Freund's adjuvant . A booster dose of 100 μ g in incomplete Freund's adjuvant was given four weeks later. Venous blood samples were obtained once a week for six weeks and were tested for reactivity for the TTV peptide 35 (SEQ ID NO:1).

Results

Human reactivities to the 97 peptides covering ORF1 and ORF2 have been given in Tables 2 and 3. Reactive peptides within ORF1 were found to be peptides 10 (SEQ ID NO:3), 18 (SEQ ID NO:4), 29 (SEQ ID NO:5), 35 (SEQ ID NO:1), 42 (SEQ ID NO:6), 44 (SEQ ID NO:7), 50 (SEQ ID NO:8), 51 (SEQ ID NO:9), and 69 (SEQ ID NO:10) (Table 2). Two of the tested human sera were reactive with peptide 19 (SEQ ID NO:11) from ORF2 (Table 3). All reactive peptides have been listed in Table 4. The most often detected peptide was the peptide 35 with the sequence TATTTTYAYPGTNRPPV (SEQ ID NO:1). The reactivity to peptide 35 was dependent on the dilution of the serum samples (Table 5). The reactivity of the human serum samples to the peptide on the microplate could be inhibited by the addition of the same peptide in solution, but not by an irrelevant peptide (data not shown). This shows that the reactivity is specific for the peptide 35 with the sequence TATTTTYAYPGTNRPPV (SEQ ID NO:1).

The reactivity to the TATTTTYAYPGTNRPPV peptide was further characterized using deletion and substitution peptide analogues. This analysis showed that the recognized region contained the sequence YAYPGTNRPPV (SEQ ID NO:2) (Table 6). Using alanine substitution analogues the Pro-Val sequence was found to be the one most essential for the binding of human antibodies (Table 6).

Table 1 Complete amino acid sequences of the ORFs 1 and 2 of TTV (Genebank accession no AB008394) used for the synthesis of 80 overlapping peptides.

ORF1

5 MAYGWRRRRRRWRRWRRRPWRRRWTRRRRPARRRGRRRNVRRRRRGGRWRR
RYRRWKRKGRRRKKAKIIRQWQPNYRRRCNIVGYIPVLICGENTVSRNYATHSDDT
NYPGPFGGGMTTDKFTLRILYDEYKRFMNYWTASNEDDLDCRYLGVNLYFFRHPDV
DFIIKINTMPPFLDTELAPSIHPGMLALDKRARWIPSLKSRPGKKHYIKIRVGAPRMFT
DKWYPQTDLCDMVLLTVYATAADMQYPFGSPLTDSVVVNFQVLQSMYDKTISILPD
10 EKSQREILLNKIASYPFYNTTQTIAQLKPFIDAGNVTS GATATTWASYINTTKFTTATT
TTYAYPGTNRPPVTMLTCNDSWYRGTVYNTQIQQLPIKAAKLYLEATKTLGNTFTN
EDYTLEYHGGLYSSIWLSPGRSYFETT GAYTDIKYNPFTDRGEGNMLWIDWLSKKN
MNYDKVQSKCLISDLPLWAAAYGYVEFCAKSTGDQNIHMNARLLIRSPFTDPQLLVH
TDPTKGFVPYSLNFGNGKMPGGSSNVPIRMRAKWYPTLFHQQEVLEALA QSGPFAY
15 HSDIKKVS LGMKYRFKWIWGGNPVRQQVVRNPCKETHSSGNRVPRSLQIVDPKYNS
PELTFHTWDFRRGLFGPKAIQRMQQPTTTDIFSAGRKRPRRDTEVYHSSQEGEQKES
LLFPPVKLLRRVPPWEDSQQEESGSQSSEETQTVSQQLKQQLQQQRILGVKLRLLFN
QVQKIQQNQDINPTLLPRGGDLASLFQIAP

ORF2

20 MAEFSTPVRSGEATEGDLRVPRAGAEGEFTHRSQGAIRARDWPGYGQGSEKSMFIGR
HYRKKRALSLCAVRTTKKACKLLIVMWTPPRNDQHLYLNWQWYSSILSSHAAMCGC
PDAVAHFNHLASVLRAPQNPPPPGPQRNLPLRRLPALPAAPEAPGDRAPWPMAGGAE
GEDGGAGGDADHGGAAGGPEDADLLDAVAAAE

25

TABLE 2. Analysis of human antibody reactivities in EIA to overlapping synthetic peptides corresponding to the complete open reading frame 1 (ORF1) of the TT virus. Values have been given as the optical density at 405 nm. OD values over 0.500 are considered positive and have been written in bold.

TTV ORF1 peptide	Samples negative for TTV DNA by PCR										Samples positive for TTV DNA by PCR									
	1	2	3	4	5	6	7	11	9	18	36	48	98	110	155	157				
1	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00				
2	0.08	0.08	0.07	0.11	0.07	0.09	0.08	0.14	0.07	0.06	0.08	0.10	0.05	0.08	0.07	0.08				
3	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00				
4	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00				
5	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00				
6	0.08	0.08	0.07	0.10	0.07	0.09	0.07	0.11	0.08	0.06	0.08	0.09	0.05	0.07	0.08	0.07				
7	0.07	0.07	0.08	0.10	0.07	0.08	0.08	0.10	0.07	0.064	0.07	0.09	0.05	0.07	0.08	0.08				
8	0.07	0.06	0.07	0.08	0.07	0.09	0.07	0.10	0.07	0.07	0.08	0.10	0.05	0.07	0.09	0.09				
9	0.07	0.08	0.08	0.09	0.08	0.10	0.07	0.10	0.08	0.06	0.07	0.10	0.05	0.08	0.10	0.11				
10	0.09	0.10	0.11	0.11	0.11	0.13	1.09	0.12	0.09	0.07	0.07	0.11	0.06	0.09	0.12	0.12				
11	0.09	0.10	0.10	0.10	0.09	0.11	0.09	0.12	0.09	0.07	0.08	0.10	0.07	0.08	0.12	0.13				
12	0.09	0.09	0.10	0.10	0.08	0.11	0.09	0.12	0.10	0.08	0.08	0.11	0.06	0.08	0.13	0.13				
13	0.09	0.08	0.08	0.09	0.08	0.09	0.09	0.10	0.09	0.07	0.08	0.10	0.07	0.08	0.11	0.11				
14	0.09	0.10	0.11	0.11	0.08	0.12	0.09	0.12	0.08	0.07	0.07	0.09	0.05	0.07	0.14	0.14				
15	0.08	0.09	0.11	0.10	0.09	0.13	0.09	0.13	0.09	0.06	0.07	0.13	0.05	0.08	0.15	0.16				
16	0.08	0.09	0.08	0.10	0.08	0.14	0.09	0.11	0.08	0.07	0.07	0.09	0.04	0.06	0.12	0.12				
17	0.08	0.10	0.09	0.11	0.09	0.13	0.09	0.13	0.08	0.08	0.08	0.11	0.04	0.09	0.12	0.13				
18	0.09	0.09	0.15	0.29	0.07	0.13	0.62	0.28	0.08	0.07	0.13	0.12	0.04	0.08	0.19	0.18				
19	0.08	0.09	0.10	0.10	0.09	0.16	0.12	0.13	0.08	0.07	0.07	0.14	0.04	0.08	0.15	0.14				
20	0.07	0.06	0.06	0.07	0.06	0.08	0.08	0.11	0.08	0.07	0.07	0.09	0.04	0.06	0.09	0.10				
21	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00				

22	0.07	0.07	0.07	0.08	0.08	0.09	0.09	0.09	0.11	0.08	0.07	0.08	0.10	0.04	0.07	0.11	0.11
23	0.09	0.09	0.11	0.11	0.08	0.16	0.09	0.12	0.12	0.08	0.07	0.07	0.10	0.05	0.07	0.14	0.14
24	0.07	0.09	0.12	0.12	0.08	0.12	0.08	0.11	0.08	0.07	0.07	0.07	0.10	0.05	0.10	0.13	0.15
25	0.07	0.09	0.08	0.08	0.06	0.11	0.07	0.12	0.12	0.07	0.06	0.06	0.06	0.06	0.07	0.11	0.12
26	0.07	0.09	0.07	0.08	0.08	0.10	0.07	0.10	0.10	0.07	0.07	0.06	0.05	0.07	0.08	0.09	0.11
27	0.07	0.08	0.07	0.08	0.06	0.11	0.07	0.08	0.08	0.06	0.06	0.06	0.06	0.08	0.06	0.09	0.10
28	0.07	0.08	0.08	0.08	0.07	0.50	0.07	0.10	0.10	0.07	0.06	0.06	0.06	0.07	0.06	0.09	0.10
29	0.42	0.36	0.45	0.53	0.22	0.95	0.29	0.14	0.21	0.11	0.09	0.09	0.10	0.09	0.25	0.59	0.62
30	0.07	0.07	0.06	0.07	0.06	0.09	0.07	0.09	0.07	0.07	0.07	0.07	0.07	0.08	0.08	0.09	0.12
31	0.05	0.09	0.09	0.08	0.06	0.09	0.07	0.08	0.06	0.06	0.06	0.06	0.06	0.06	0.06	0.09	0.09
32	0.06	0.08	0.06	0.25	0.06	0.09	0.06	0.07	0.06	0.05	0.05	0.06	0.05	0.07	0.06	0.09	0.09
33	0.07	0.08	0.06	0.07	0.06	0.08	0.06	0.07	0.06	0.05	0.05	0.06	0.05	0.07	0.06	0.08	0.09
34	0.07	0.07	0.05	0.06	0.06	0.06	0.06	0.07	0.06	0.09	0.09	0.06	0.15	0.06	0.06	0.08	0.08
35	0.18	0.17	0.27	1.40	0.25	0.64	2.10	0.64	0.10	0.10	0.10	1.22	1.11	0.05	0.47	0.95	0.86
36	0.07	0.07	0.06	0.07	0.06	0.11	0.07	0.08	0.09	0.06	0.06	0.06	0.07	0.07	0.07	0.08	0.08
37	0.10	0.11	0.11	0.12	0.09	0.14	0.12	0.14	0.09	0.08	0.08	0.08	0.20	0.10	0.08	0.14	0.14
38	0.07	0.08	0.08	0.09	0.08	0.11	0.09	0.11	0.08	0.07	0.07	0.07	0.07	0.09	0.08	0.19	0.11
39	0.08	0.09	0.07	0.09	0.07	0.37	0.09	0.10	0.08	0.09	0.09	0.09	0.08	0.09	0.09	0.11	0.13
40	0.09	0.16	0.24	0.12	0.11	0.16	0.09	0.12	0.09	0.09	0.09	0.08	0.08	0.10	0.09	0.15	0.16
41	0.08	0.12	0.10	0.11	0.10	0.20	0.09	0.15	0.09	0.07	0.07	0.09	0.08	0.11	0.09	0.15	0.16
42	0.08	0.11	0.09	0.10	0.17	0.23	0.11	0.19	0.27	0.12	0.12	0.10	0.09	0.16	0.42	0.35	1.24
43	0.09	0.21	0.11	0.11	0.10	0.41	0.10	0.14	0.08	0.08	0.08	0.08	0.08	0.12	0.11	0.20	0.25
44	0.09	0.11	0.11	0.12	0.98	0.15	0.10	0.14	0.09	0.09	0.09	0.08	0.08	0.12	0.13	0.17	1.25
45	0.08	0.06	0.06	0.12	0.10	0.17	0.10	0.14	0.09	0.08	0.08	0.06	0.09	0.11	0.09	0.16	0.17
46	0.08	0.08	0.07	0.09	0.08	0.22	0.10	0.12	0.08	0.07	0.07	0.07	0.07	0.09	0.08	0.19	0.26
47	0.07	0.09	0.08	0.09	0.09	0.13	0.09	0.12	0.07	0.07	0.07	0.07	0.07	0.10	0.09	0.15	0.19
48	0.08	0.11	0.10	0.12	0.10	0.16	0.11	0.13	0.08	0.09	0.09	0.08	0.08	0.12	0.09	0.15	0.15
49	0.09	0.11	0.10	0.11	0.10	0.15	0.15	0.15	0.08	0.08	0.08	0.07	0.08	0.10	0.11	0.26	0.25
50	0.07	0.09	0.08	0.09	0.08	0.86	0.29	0.18	0.08	0.07	0.07	0.07	0.07	0.10	0.20	0.63	0.46

51	0.06	0.09	0.08	0.10	0.12	0.24	0.13	0.13	0.07	0.07	0.08	0.08	0.11	0.09	0.24	0.78
52	0.10	0.13	0.11	0.11	0.10	0.20	0.11	0.13	0.10	0.09	0.08	0.08	0.14	0.11	0.21	0.38
53	0.12	0.12	0.09	0.10	0.08	0.16	0.09	0.11	0.09	0.10	0.08	0.07	0.09	0.10	0.16	0.15
54	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
55	0.10	0.10	0.07	0.08	0.07	0.14	0.09	0.08	0.35	0.08	0.07	0.07	0.08	0.09	0.14	0.12
56	0.10	0.16	0.11	0.11	0.09	0.18	0.09	0.11	0.10	0.09	0.09	0.09	0.10	0.10	0.18	0.15
57	0.14	0.12	0.10	0.12	0.09	0.17	0.11	0.12	0.11	0.08	0.09	0.09	0.10	0.12	0.19	0.43
58	0.07	0.06	0.06	0.07	0.06	0.07	0.07	0.08	0.07	0.06	0.08	0.08	0.04	0.06	0.08	0.09
59	0.06	0.05	0.05	0.06	0.06	0.06	0.06	0.08	0.06	0.05	0.05	0.06	0.04	0.05	0.07	0.08
60	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
61	0.07	0.06	0.06	0.08	0.06	0.10	0.07	0.10	0.08	0.05	0.06	0.08	0.04	0.07	0.12	0.12
62	0.07	0.07	0.06	0.09	0.07	0.08	0.07	0.09	0.08	0.05	0.07	0.11	0.05	0.07	0.11	0.14
63	0.11	0.16	0.12	0.13	0.11	0.16	0.11	0.05	0.08	0.08	0.08	0.08	0.11	0.09	0.15	0.15
64	0.09	0.08	0.08	0.09	0.07	0.10	0.08	0.09	0.08	0.08	0.08	0.07	0.08	0.08	0.11	0.12
65	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
66	0.10	0.12	0.10	0.10	0.10	0.14	0.12	0.16	0.10	0.09	0.09	0.09	0.11	0.10	0.13	0.15
67	0.10	0.11	0.09	0.12	0.10	0.15	0.12	0.15	0.10	0.09	0.09	0.09	0.10	0.10	0.21	0.24
68	0.11	0.14	0.16	0.16	0.13	0.23	0.13	0.19	0.10	0.10	0.09	0.10	0.12	0.12	0.20	0.21
69	0.10	0.15	0.16	0.14	0.12	0.51	0.11	0.13	0.09	0.08	0.08	0.09	0.10	0.11	0.15	0.21
70	0.09	0.11	0.11	0.11	0.09	0.19	0.12	0.12	0.08	0.08	0.08	0.09	0.10	0.11	0.18	0.20
71	0.10	0.13	0.14	0.14	0.11	0.20	0.11	0.17	0.10	0.08	0.10	0.09	0.13	0.14	0.21	0.20
72	0.10	0.15	0.12	0.13	0.10	0.19	0.12	0.16	0.08	0.08	0.09	0.09	0.11	0.11	0.18	0.19
73	0.08	0.09	0.09	0.10	0.09	0.12	0.10	0.13	0.09	0.08	0.07	0.08	0.11	0.09	0.13	0.14
74	0.08	0.09	0.07	0.08	0.08	0.10	0.09	0.11	0.09	0.08	0.07	0.08	0.08	0.08	0.09	0.11
75	0.11	0.12	0.12	0.11	0.10	0.17	0.44	0.15	0.10	0.10	0.10	0.09	0.11	0.10	0.15	0.16
76	0.10	0.13	0.15	0.14	0.11	0.28	0.13	0.14	0.11	0.09	0.08	0.09	0.12	0.26	0.01	0.22
77	0.14	0.14	0.15	0.12	0.09	0.18	0.12	0.13	0.18	0.07	0.07	0.08	0.08	0.09	0.11	0.14

Table 3. Analysis of human antibody reactivities in EIA to overlapping synthetic peptides corresponding to the complete open reading frame 2 (ORF2) of the TT virus. Values have been given as the optical density (OD) at 405 nm. OD values over 0.500 are considered positive and have been written in bold.

TTV ORF2 peptide	Samples negative for TTV DNA by PCR										Samples positive for TTV DNA by PCR									
	1	2	3	4	5	6	7	11	9	18	36	48	98	110	155	157				
1	0.09	0.10	0.13	0.11	0.10	0.16	0.09	0.12	0.08	0.08	0.09	0.08	0.08	0.08	0.14	0.14				
2	0.10	0.11	0.11	0.10	0.08	0.12	0.09	0.14	0.08	0.08	0.08	0.08	0.08	0.09	0.14	0.13				
3	0.08	0.10	0.11	0.10	0.08	0.29	0.28	0.14	0.10	0.08	0.13	0.08	0.11	0.10	0.14	0.04				
4	0.09	0.09	0.08	0.08	0.07	0.09	0.07	0.09	0.08	0.07	0.07	0.07	0.07	0.07	0.10	0.10				
5	0.08	0.08	0.08	0.08	0.07	0.10	0.08	0.10	0.07	0.07	0.07	0.07	0.07	0.08	0.12	0.13				
6	0.07	0.10	0.08	0.10	0.07	0.10	0.07	0.10	0.08	0.28	0.09	0.08	0.10	0.09	0.09	0.11				
7	0.11	0.10	0.11	0.14	0.91	0.13	0.10	0.11	0.08	0.09	0.10	0.11	0.10	0.09	0.10	0.13				
8	0.09	0.10	0.11	0.13	0.10	0.12	0.10	0.14	0.09	0.08	0.10	0.11	0.10	0.10	0.11	0.12				
9	0.08	0.09	0.09	0.09	0.09	0.10	0.09	0.11	0.09	0.08	0.08	0.09	0.09	0.09	0.13	0.14				
10	0.09	0.11	0.10	0.09	0.08	0.11	0.08	0.10	0.09	0.09	0.07	0.08	0.08	0.08	0.11	0.13				
11	0.10	0.10	0.11	0.08	0.09	0.09	0.09	0.11	0.10	0.12	0.09	0.07	0.09	0.09	0.14	0.14				
12	0.12	0.09	0.09	0.08	0.09	0.09	0.08	0.07	0.13	0.09	0.08	0.10	0.14	0.12	0.16	0.14				
13	0.07	0.09	0.09	0.08	0.08	0.10	0.09	0.11	0.07	0.08	0.07	0.08	0.09	0.08	0.12	0.12				
14	0.08	0.07	0.08	0.07	0.07	0.08	0.08	0.10	0.07	0.07	0.07	0.07	0.07	0.07	0.10	0.10				
15	0.08	0.08	0.08	0.07	0.07	0.09	0.07	0.10	0.08	0.07	0.07	0.07	0.08	0.08	0.10	0.12				
16	0.08	0.09	0.09	0.07	0.07	0.09	0.08	0.09	0.08	0.08	0.08	0.07	0.08	0.08	0.12	0.11				
17	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00				
18	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00				
19	0.09	0.12	0.17	0.12	0.08	0.51	0.57	0.15	0.08	0.07	0.33	0.07	0.08	0.08	0.14	0.14				
20	0.09	0.10	0.09	0.08	0.07	0.10	0.07	0.09	0.10	0.10	0.08	0.08	0.08	0.10	0.14	0.16				

TABLE 4. Sequences of TTV peptides reactive with human serum samples.**ORF1****Peptide no. Peptide sequence**

10	VLICGENTVSRNYATHS	SEQ ID NO:3
18	KINTMPPFLDTELTAPS	SEQ ID NO:4
29	PDEKSQREILLNKIASY	SEQ ID NO:5
35	TATTTTYAYPGTNRPPV	SEQ ID NO:1
42	GLYSSIWLSPGRSYFET	SEQ ID NO:6
44	YTDIKYNPFTDRGEGNM	SEQ ID NO:7
50	DQNIHMNARLLIRSPFT	SEQ ID NO:8
51	LIRSPFTDPQLLVHTDP	SEQ ID NO:9
69	QKESLLFPPVKLLRRVP	SEQ ID NO:10

ORF2**Peptide no. Peptide sequence**

19	EDGGAGGDADHGGAAGGP	SEQ ID NO:11
----	--------------------	--------------

TABLE 5. Analysis of the reactivities of serial dilutions of three human serum samples with to the TTV peptide TATTTTYAYPGTNRPPV (SEQ ID NO:1). Values are given as the OD and standard deviation (SD) at 405 nm.

Dilution of serum sample	Human serum sample					
	P 4	SD	P 6	SD	P 7	SD
1:100	1.285	0.072	0.687	0.082	1.782	0.054
1:200	0.758	0.056	0.375	0.003	1.23	0.02
1:400	0.411	0.021	0.19	0.007	0.79	0.018
1:800	0.234	0.008	0.104	0.003	0.45	0.002
1:1600	0.131	0.005	0.067	0.001	0.246	0.013
1:3200	0.076	0.003	0.049		0.131	0.006
1:6400	0.058	0.001	0.043		0.087	
1:12800	0.046		0.041		0.061	

TABLE 6. Analysis of the reactivities of three human serum samples with to the deletion and alanine substitution analogues of the TTV peptide TATTTTYAYPGTNRPPV (SEQ ID NO:1). Values are given as the OD at 405 nm. Positive reactivities, i.e. more than 50% of the reactivity of the original peptide, have been written in bold.

Deletion or substitution peptide analogue	Human serum sample		
	P4	P7	P36
TATTTTYAYPGTNRPPV	0.839	1.845	0.825
TATTTTYAYPGTNRPP	0.096	0.144	0.086
TATTTTYAYPGTNRP	0.100	0.099	0.078
TATTTTYAYPGTNR	0.092	0.103	0.078
TATTTTYAYPGTN	0.186	0.888	0.083
TATTTTYAYPGT	0.095	0.087	0.072
TATTTTYAYPG	0.095	0.085	0.074
TATTTTYAYP	0.095	0.096	0.082
TATTTTYAY	0.098	0.089	0.083
TATTTTYA	0.115	0.089	0.090
TATTTTY	0.142	0.105	0.076
TATTTT	0.108	0.093	0.082
TATTT	0.101	0.091	0.078
TATT	0.099	0.105	0.076
ATTTTYAYPGTNRPPV	1.042	1.960	0.923
TTTTYAYPGTNRPPV	0.805	1.587	0.776
TTYAYPGTNRPPV	0.697	1.488	0.810
TYAYPGTNRPPV	0.748	1.659	0.722
YAYPGTNRPPV	0.707	1.508	0.712
AYPGTNRPPV	0.647	1.546	0.677
YPGTNRPPV	0.662	1.488	0.669
PGTNRPPV	0.300	1.091	0.406
GTNRPPV	0.166	0.430	0.123
TNRPPV	0.300	0.887	0.210
NRPPV	0.110	0.146	0.056
AAATTTTYAYPGTNRPPV	0.135	0.242	0.076
TGTTTTYAYPGTNRPPV	1.045	1.852	0.915
TAATTTTYAYPGTNRPPV	0.855	1.829	0.806
TATATTTTYAYPGTNRPPV	0.897	1.675	0.764
TATTATYAYPGTNRPPV	0.971	1.722	0.824
TATTTAYAYPGTNRPPV	1.076	1.867	0.955
TATTTTAAAYPGTNRPPV	1.011	1.833	1.027
TATTTTGYPGTNRPPV	0.898	1.619	0.901
TATTTTYAGTNRPPV	0.836	1.769	0.850
TATTTTYAAPGTNRPPV	0.899	1.697	0.903
TATTTTYAYAGTNRPPV	0.886	1.738	0.903
TATTTTYAYPATNRPPV	0.895	1.503	0.734
TATTTTYAYPGANRPPV	0.891	1.594	0.714
TATTTTYAYPGTARPPV	1.226	1.723	0.696
TATTTTYAYPGTNAPPV	0.761	1.558	0.708
TATTTTYAYPGTNRAPV	0.720	1.551	0.812
TATTTTYAYPGTNRPAV	0.090	0.092	0.100
TATTTTYAYPGTNRPPA	0.108	0.105	0.095

References

1. Nishizawa T, Okamoto H, Konishi K, Yoshizawa H, Miyakawa Y, Mayumi M. A novel DNA virus (TTV) associated with elevated transaminase levels in posttransfusion hepatitis of unknown etiology. *Biochem Biophys Res Commun* 1997;241:92-7
2. Okamoto H, Akahane Y, Ukita M, Fukuda M, Tsuda F, Miyakawa Y, Mayumi M. Fecal excretion of a nonenveloped DNA virus (TTV) associated with posttransfusion non-A-G hepatitis. *J Med Virol* 1998;56:128-32
3. Cossart Y. TTV a common virus, but pathogenic? [comment]. *Lancet* 1998;352:164
4. Okamoto H, Kato N, Iizuka H, Tsuda F, Miyakawa Y, Mayumi M. Distinct genotypes of a nonenveloped DNA virus associated with posttransfusion non-A to G hepatitis (TT virus) in plasma and peripheral blood mononuclear cells [In Process Citation]. *J Med Virol* 1999;57:252-8
5. Naoumov NV, Petrova EP, Thomas MG, Williams R. Presence of a newly described human DNA virus (TTV) in patients with liver disease [see comments]. *Lancet* 1998;352:195-7
6. Viazov S, Ross RS, Varenholz C, Lange R, Holtmann M, Niel C, Roggendorf M. Lack of evidence for an association between TTV infection and severe liver disease [In Process Citation]. *J Clin Virol* 1998;11:183-7
7. Sällberg M, Ruden U, Magnus LO, Norrby E, Wahren B. Rapid "tea-bag" peptide synthesis using 9-fluorenylmethoxycarbonyl (Fmoc) protected amino acids applied for antigenic mapping of viral proteins. *Immunology Letters* 1991;30:59-68
8. Zhang ZX, Chen M, Hultgren C, Birkett A, Milich DR, Sällberg M. Immune responses to the hepatitis C virus NS4a are profoundly influenced by the combination of the viral genotype and the host major histocompatibility complex. *J. Gen. Virol.* 1997;78:2735-2746

CLAIMS

1. Peptide having the amino acid sequence

SEQ ID NO:1

- 5 Thr Ala Thr Thr Thr Thr Tyr Ala Tyr Pro Gly Thr Asn Arg Pro Pro
Val

wherein one to six of the N-terminal amino acids Thr Ala Thr Thr Thr Thr may be omitted.

2. Peptide according to claim 1 having the amino acid sequence

SEQ ID NO:2

- 10 Tyr Ala Tyr Pro Gly Thr Asn Arg Pro Pro Val.

3. Peptide mixture comprising the peptide according to claim 1 and at least one of the peptides

SEQ ID NO:3

Val Leu Ile Cys Gly Glu Asn Thr Val Ser Arg Asn Tyr Ala Thr His

- 15 Ser,

SEQ ID NO:4

Lys Ile Asn Thr Met Pro Pro Phe Leu Asp Thr Glu Leu Thr Ala Pro
Ser,

SEQ ID NO:5

- 20 Pro Asp Glu Lys Ser Gln Arg Glu Ile Leu Leu Asn Lys Ile Ala Ser
Tyr,

SEQ ID NO:6

Gly Leu Tyr Ser Ser Ile Trp Leu Ser Pro Gly Arg Ser Tyr Phe Glu
Thr,

- 25 SEQ ID NO:7

Tyr Thr Asp Ile Lys Tyr Asn Pro Phe Thr Asp Arg Gly Glu Gly Asn
Met,

SEQ ID NO:8

Asp Gln Asn Ile His Met Asn Ala Arg Leu Leu Ile Arg Ser Pro Phe
30 Thr,

SEQ ID NO:9

Leu Ile Arg Ser Pro Phe Thr Asp Pro Gln Leu Leu Val His Thr Asp
Pro,

SEQ ID NO:10

Gln Lys Glu Ser Leu Leu Phe Pro Pro Val Lys Leu Leu Arg Arg Val

Pro, and

SEQ ID NO:11

5 Glu Asp Gly Gly Ala Gly Gly Asp Ala Asp His Gly Gly Ala Ala Gly
Gly Pro.

4. Peptide according to claim 1 or 2, or a peptide mixture according to claim 3,
wherein at least one peptide is coupled to a carrier and/or label.

5. Peptide according to claim 1 or 2, or a peptide mixture according to claim 3,
10 wherein at least one peptide is immobilized on a solid phase.

6. Peptide or peptide mixture according to any one of the preceding claims for use in
a medicament.

7. Monospecific antibody binding to the TT virus,

8. Monospecific antibody according to claim 7 binding to an amino acid sequence
15 selected from the group consisting of the amino acid sequences SEQ ID NO:1 - 11.

9. Monospecific antibody according to claim 7 or 8 for use in a medicament.

10. Diagnostic kit comprising a peptide or peptide mixture according to any one of
claims 1 - 5 as diagnostic antigen(s).

11. Diagnostic kit comprising one or more monospecific antibodies according to claim
20 7 or 8 as diagnostic antibodies.

12. Use of a peptide or peptide mixture according to any one of claims 1 - 4 for
immunization of a non-human mammal to produce monospecific antibodies directed against
TT virus.

SEQUENCE LISTING

<110> Tripep AB

<120> Peptides from the TT virus sequence and monospecific
antibodies binding to the TT virus

<130> 192975901

<140>

<141>

<160> 11

<170> PatentIn Ver. 2.1

<210> 1

<211> 17

<212> PRT

<213> TT virus

<400> 1

Thr Ala Thr Thr Thr Tyr Ala Tyr Pro Gly Thr Asn Arg Pro Pro
1 5 10 15

Val

<210> 2

<211> 11

<212> PRT

<213> TT virus

<400> 2

Tyr Ala Tyr Pro Gly Thr Asn Arg Pro Pro Val
1 5 10

<210> 3

<211> 17

<212> PRT

<213> TT virus

<400> 3

Val Leu Ile Cys Gly Glu Asn Thr Val Ser Arg Asn Tyr Ala Thr His
1 5 10 15

Ser

<210> 4

<211> 17

<212> PRT

<213> TT virus

<400> 4

Lys Ile Asn Thr Met Pro Pro Phe Leu Asp Thr Glu Leu Thr Ala Pro
1 5 10 15

Ser

<210> 5
<211> 17
<212> PRT
<213> TT virus

<400> 5
Pro Asp Glu Lys Ser Gln Arg Glu Ile Leu Leu Asn Lys Ile Ala Ser
1 5 10 15

Tyr

<210> 6
<211> 17
<212> PRT
<213> TT virus

<400> 6
Gly Leu Tyr Ser Ser Ile Trp Leu Ser Pro Gly Arg Ser Tyr Phe Glu
1 5 10 15

Thr

<210> 7
<211> 17
<212> PRT
<213> TT virus

<400> 7
Tyr Thr Asp Ile Lys Tyr Asn Pro Phe Thr Asp Arg Gly Glu Gly Asn
1 5 10 15

Met

<210> 8
<211> 17
<212> PRT
<213> TT virus

<400> 8
Asp Gln Asn Ile His Met Asn Ala Arg Leu Leu Ile Arg Ser Pro Phe
1 5 10 15

Thr

<210> 9
<211> 17
<212> PRT
<213> TT virus

<400> 9
Leu Ile Arg Ser Pro Phe Thr Asp Pro Gln Leu Leu Val His Thr Asp
1 5 10 15

Pro

<210> 10
<211> 17
<212> PRT
<213> TT virus

<400> 10
Gln Lys Glu Ser Leu Leu Phe Pro Pro Val Lys Leu Leu Arg Arg Val
1 5 10 15

Pro

<210> 11
<211> 18
<212> PRT
<213> TT virus

<400> 11
Glu Asp Gly Gly Ala Gly Gly Asp Ala Asp His Gly Gly Ala Ala Gly
1 5 10 15

Gly Pro

INTERNATIONAL SEARCH REPORT

International Application No
PCT/EP 00/03958

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07K14/01 C07K16/08 G01N33/68 A61P31/20

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07K G01N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ, MEDLINE, BIOSIS, STRAND

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P, A	WO 99 58638 A (INNOGENETICS NV ; MAERTENS GEERT (BE); VREESE KAREN DE (BE)) 18 November 1999 (1999-11-18) claims 21-38	1,6-12
A	WO 99 05282 A (NISHIZAWA TSUTOMU ; OKAMOTO HIROAKI (JP); TAMURA RYOJI (JP)) 4 February 1999 (1999-02-04) claims; examples - & EP 1 010 759 A 21 June 2000 (2000-06-21) -/--	1,6-12

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

4 September 2000

Date of mailing of the international search report

13/09/2000

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Fuhr, C

INTERNATIONAL SEARCH REPORT

International Application No
PCT/EP 00/03958

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
T	<p>DATABASE WPI Section Ch, Week 200036 Derwent Publications Ltd., London, GB; Class B04, AN 2000-415430 XP002146537 & JP 2000 135087 A (SRL KK), 16 May 2000 (2000-05-16) abstract</p> <p>-----</p>	1,6-12

INTERNATIONAL SEARCH REPORT

Information on patent family members

Inte. Appl. No.

PCT/EP 00/03958

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9958638 A	18-11-1999	AU 4259299 A	29-11-1999
WO 9905282 A	04-02-1999	AU 8358698 A	16-02-1999
		EP 1010759 A	21-06-2000
		ZA 9809239 A	19-04-1999
JP 2000135087 A	16-05-2000	NONE	